Health Certificate No
(Valid Only if the USDA Veterinary
Seal Appears Over the Certificate #)

"The list of team veterinarians approved to export bovine embryos to the European Union is listed in the IRRS file EE.EM.VT."

October 2002

## HEALTH CERTIFICATE FOR THE EXPORTATION OF BOVINE EMBRYOS TO THE EUROPEAN COMMUNITY (EC)

1.	Cons	signor (name and full address)	NoORIGINAL		
			2.	Third Country of Collection	
3.	Cons	signee (name and full address)	4.	Competent Authority	
NOT	ES: (a)	A separate certificate must	5.	Competent Local Authority	
	(b)	be issued for each consignment of embryos The original of this certificate must accompany the consignment			
6.	Place	e and date of loading	7.	Name and address of embryo collection team or embryo production team(1)	
8.	Mean	ns of transport	9.	Place and Member State of destination	
10.	_	stration number of embryo collection or embryo production team (1)	11.	Number and codemark of embryo containers	

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Embryos (a)	Identification of consignment:  Embryos (a) derived by in vitro fertilization yes/no (1)  (b) subjected to penetration of zona pellucida yes/no (1)					
a) Number o	of embryo	os (b)	) Date(s) of collection	(c) Breed		
I, the undersi certify that:	gned offi	cial veter	inarian of the Governmen	nt of <u>the United States</u> (name of exporting third country)		
1.	The em	ibryo coll	ection/production team ic	dentified above:		
		proved in 56/EEC.	accordance with Chapter	1 of Annex A to Directive		
	transp	ort of the		production and storing and e in accordance with Chapter II		
		ojected at inarian;	least twice per year to in	spection by an official		
2.		ling to of	ficial findings the United ry) has:	States (name of		
	(a)	been free during 12 months immediately prior to collection o embryos to be exported from rinderpest:				
	(b)	-	):			
		n	nonths immediately prior of be exported and does no	d-mouth disease during the 12 to the collection of the embryos of practice vaccination against it		
		(ii) h	as not been free from foo nonths immediately prior nd/or practices vaccination the donor females a and other tissues us come from a holdir	and donors of ovaries, oocytes sed in the production of embryos ag in which no animal has been foot-and mouth disease during		

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the embryos have been stored in approved conditions for a minimum period of 30 days immediately after collection;

## (c) either (1):

- (i) has been free from bluetongue and epizootic haemorrhagic disease (EHD) for the 12 months immediately prior to collection of the embryos to be exported and does not practice vaccination against them
- (ii) has not been free from bluetongue and epizootic haemorrhagic disease (EHD) for the 12 months immediately prior to collection of the embryos to be exported and/or practices vaccination against them and
  - the embryos have been stored in approved conditions for a minimum period of 30 days immediately after collection, and
  - the donor females and the donors of ovaries, oocytes or other tissues used in the production of embryos were subjected with negative results to an agar gel immuno diffusion test and a serum neutralization test for epizootic haemorrhagic disease antibodies on a blood sample taken not less than 21 days following collection;
- 3. (a) the premises on which the embryos to be exported or the ovaries, oocytes or other tissues used in the production of embryos to be exported were collected and processed was at the time of collection situated in the centre of an area of 20 km diameter in which according to official findings there had been no incidence of foot-and-mouth disease, bluetongue, epizootic haemorragic disease, contagious vesicular stomatitis, Rift Valley fever or contagious bovine pleuropneumonia for 30 days immediately prior to collection and in the case of embryos certified under 2(b)(ii) and (c)(ii) for 30 days after collection;
  - (b) between the time of collection or production of the embryos to be exported and their dispatch, they were stored continuously in approved premises which were situated in the center of an area of 20 km in diameter in which according to official findings there was no incidence of foot-and-mouth disease, contagious vesicular stomatitis or Rift Valley fever;

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- 4. the donor females and the donors of ovaries, oocytes or other tissues used in production of embryos:
  - (a) during the 30 days immediately prior to collection of the embryos to be exported, were located in premises situated in the center of an area of 20 km in diameter in which according to official findings there was no incidence of foot-and-mouth disease, bluetongue, epizootic haemorrhagic disease, contagious vesicular stomatitis, Rift Valley fever or contagious bovine pleuropneumonia;
  - (b) showed no clinical sign of disease on the day of collection;
  - (c) have spent the 6 months immediately prior to collection in the territory of <u>the United States</u> (name of exporting country) in a maximum of two herds which are:
    - according to official findings free from tuberculosis,
    - according to official findings free from brucellosis,
    - free from enzootic bovine leukosis or a herd or herds which has/have shown no clinical signs of enzootic bovine leukosis during the previous 3 years.
    - a herd or herds which has/have shown no clinical sign of infectious bovine rhinotracheitis/infectious pustular vulvovaginitis during the previous 12 months
- 5. the embryos to be exported were conceived as a result of artificial insemination or in vitro fertilization with semen from a donor sire standing at a semen collection centre approved by the competent authority for the collection, processing and storage of semen or with semen imported from the European Community.

Done at	
Signature(2)	
Name and qualification (in block letters):	Stamp (2)

**NOTE:** This certificate must:

- (a) be drawn up in at least the official language of the Member State of destination and the Member State where the embryos will enter Community territory;
- (b) be made out to a single consignee;
- (c) accompany the embryos in the original.

<sup>(1)</sup> Delete as appropriate.

<sup>(2)</sup> The signature and the stamp must be in a colour different to that of printing.